

Amendments to the Claims:

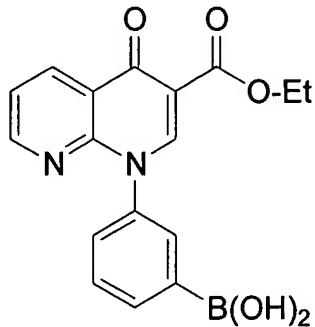
This listing of the Claims will replace all prior versions, and listings, of the claims in the application:

Listing of the Claims:

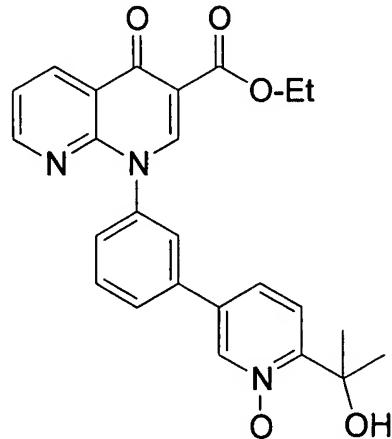
Claim 1 (Canceled)

Claim 2 (Once Amended) A method according to ~~claim 1~~ Claim 14

wherein the compound of formula Va is



and the compound of Formula VIII is



Claim 3 (Once Amended) A method according to ~~claim 1~~ Claim 14 wherein the second salt is a carbonate base.

Claim 4 (Once Amended) A method according to ~~claim 1~~ Claim 14 wherein the phosphine ligand is selected from the group consisting of P(C₁-6alkyl)₃, such as P(t-butyl)₃, P(Cy)₃, and P(t-butyl)₂(biphenyl).

Claim 5 (Once Amended) A method according to ~~claim 1~~ Claim 14 wherein the palladium catalyst is selected from the group consisting of P(t-butyl)₃-Pd-P(t-butyl)₃, [PdCl(allyl)]₂, Pd₂ (dba)₃, and [P(t-butyl)₃PdBr]₂ (Johnson-Matthey catalyst).

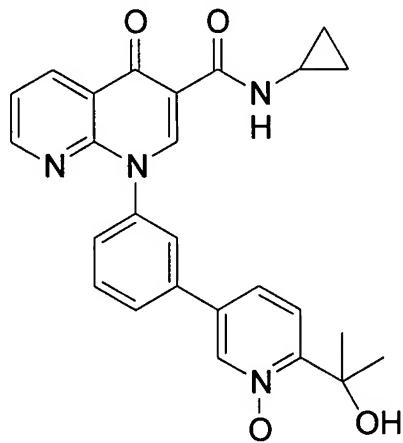
Claim 6 (Once Amended) A method according to ~~claim 1~~ Claim 14 wherein the second base is selected from sodium or potassium carbonate and sodium or potassium phosphate.

Claims 7 to 9 (Canceled)

Claim 10 (Once Amended) A method according to ~~claim 8~~ Claim 14 wherein the activating agent is selected from carbonyl diimidazole and 1-(3-Dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride.

Claims 11 to 13 (canceled)

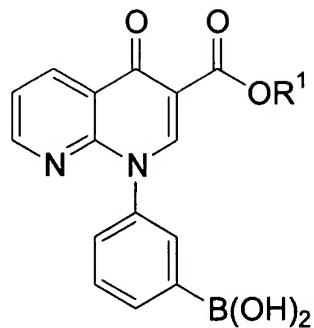
Claim 14 (New) A method of preparing a compound of Formula IX



IX

Or a pharmaceutically acceptable salt thereof, comprising

Step C: reacting, in solvent A, a compound of Formula Va

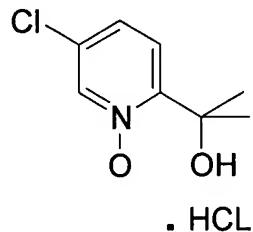


Va

wherein

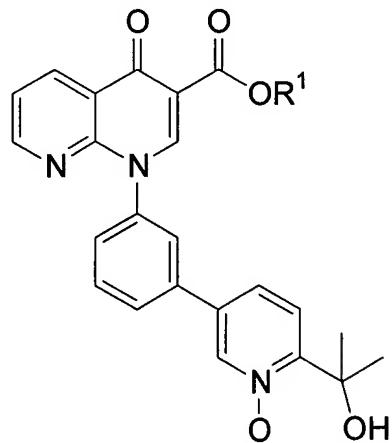
-OR¹ is a suitable leaving group; and

solvent A is selected from the group consisting of dimethylacetamide, dimethylformamide, acetonitrile, DMSO, methylacetamide, ethers or mixtures thereof; with a compound of Formula VII



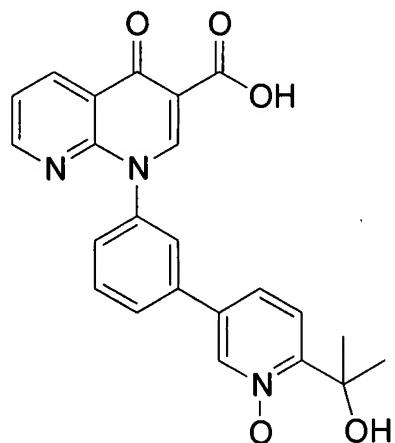
VII

or free base thereof, in the presence of a palladium catalyst and a phosphine ligand and a second base to yield a compound of Formula VIII



VIII

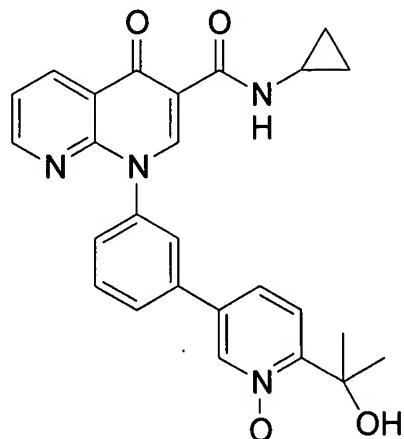
Step D: reacting, in water a compound of Formula VIII with sodium or potassium hydroxide to yield a compound of Formula VIIa



VIIa

and

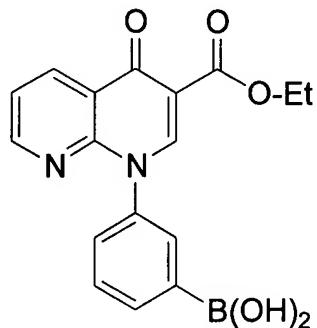
Step E: reacting, in solvent B, a compound of Formula VIIa with cyclopropylamine in the presence of an activating agent to yield a compound of Formula IX.



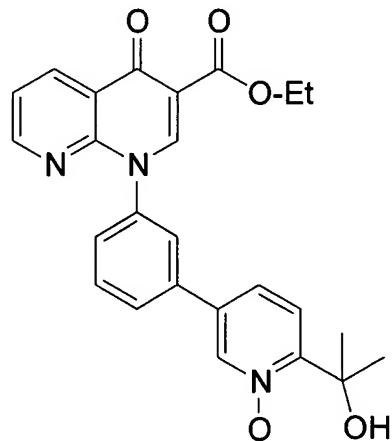
IX

wherein solvent B is selected from the group consisting of dimethylaminoacetamide, dimethylformamide, acetonitrile, DMSO, methylacetamide, dichloromethane, ethers or mixtures thereof.

Claim 15 (New) A method according to Claim 14 wherein the compound of formula Va is



and the compound of Formula VIII is



the second salt is a carbonate base,

the phosphine ligand is selected from the group consisting of P(C₁₋₆alkyl)₃, such as P(t-butyl)₃, P(Cy)₃, and P(t-butyl)₂(biphenyl),

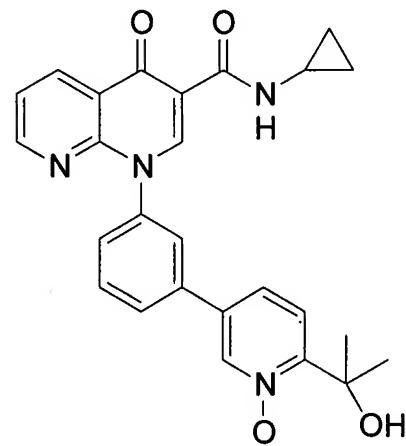
the palladium catalyst is selected from the group consisting of P(t-butyl)₃-Pd-P(t-butyl)₃,

[PdCl(allyl)]₂, Pd₂ (dba)₃, and [P(t-butyl)₃PdBr]₂ (Johnson-Matthey catalyst),

the second base is selected from sodium or potassium carbonate and sodium or potassium phosphate, and

the activating agent is selected from carbonyl diimidazole and 1-(3-Dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride.

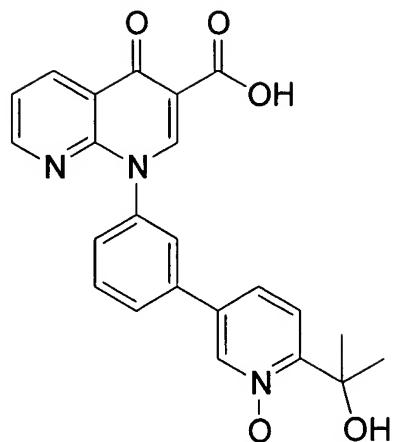
16. (New) A method of preparing a compound of Formula IX



IX

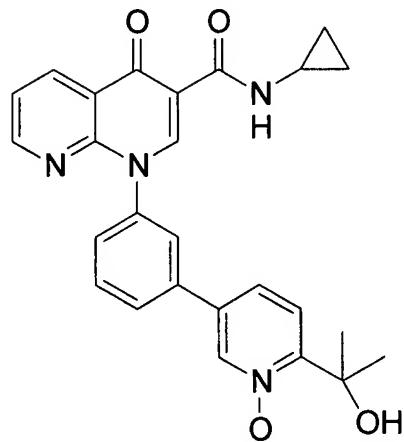
Comprising

Step E: reacting, in solvent B, a compound of Formula VIIa



VIIIa

with cyclopropylamine in the presence of an activating agent to yield a compound of Formula IX



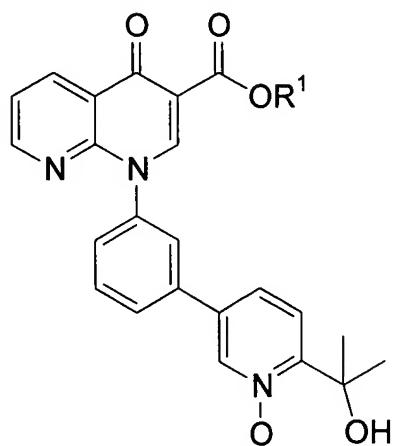
IX

wherein solvent B is selected from the group consisting of dimethylaminoacetamide, dimethylformamide, acetonitrile, DMSO, methylacetamide, dichloromethane, ethers or mixtures thereof.

17. (New) A method according to Claim 16 wherein the activating agent is selected from carbonyl diimidazole and 1-(3-Dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride.

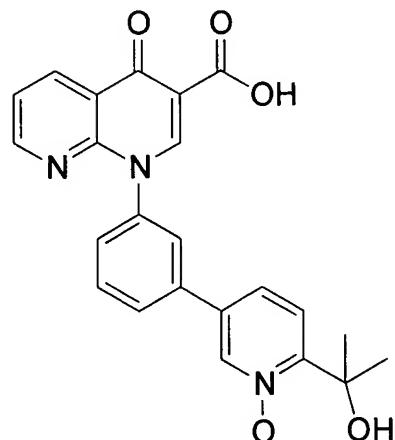
18. (New) A method according to Claim 16 further comprising

Step D: reacting, in water a compound of Formula VIII



VIII

with sodium or potassium hydroxide to yield a compound of Formula VIIIa.



VIIIa

19. (New) A method according to Claim 18 wherein the activating agent is selected from carbonyl diimidazole and 1-(3-Dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride.

20. (New) A method according to claim 18 wherein reaction step D and reaction Step E are carried out without purification or isolation of the product of Step D prior to proceeding with Step E.